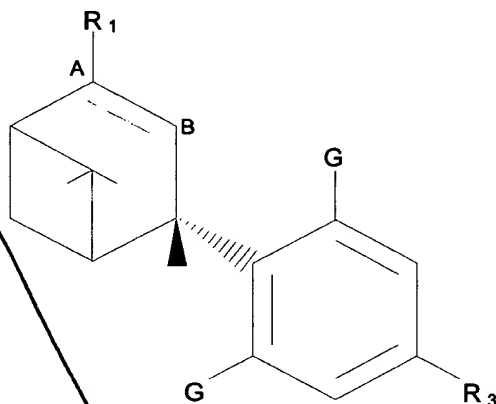


THE CLAIMS

What is claimed is:

- 5 1. A pharmaceutical composition for treating or preventing hypertension, inflammation, peripheral pain, gastrointestinal disorders, or autoimmune diseases, comprising as an active ingredient a compound of the general formula:



- 10 having the (3S,4S) configuration, and which is essentially free of the (3R,4R) enantiomer, wherein:

the dashed line A---B designates an optional double bond,

- 15 R₁ is (a) -R'N(R'')₂ wherein R' is C₁-C₅ straight or branched chain alkyl and each R'', which may be the same or different, is hydrogen or C₁-C₅ straight or branched chain alkyl optionally containing a terminal -OR''' or -OC(O)R''' moiety wherein R''' is hydrogen or C₁-C₅ straight or branched chain alkyl, (b) -Q wherein Q is a heterocyclic moiety having a labile hydrogen atom so that said moiety acts as a carboxylic acid analogue, (c) -R'X wherein R' is C₁-C₅ straight or branched chain alkyl and X is halogen, (d) -R'C(O)N(R'')₂ wherein R' is a direct bond or C₁-C₅ straight or branched chain alkyl and each R'', which may be the same or different, is hydrogen or C₁-C₅ straight or branched chain alkyl optionally containing a terminal -OR''' or -OC(O)R''' moiety wherein R''' is hydrogen or C₁-C₅ straight or branched chain alkyl, (e) -R'C(O)OR'' wherein R' is a direct bond or C₁-C₅ straight or branched chain alkyl and R'' is hydrogen or C₁-C₅ straight or branched chain alkyl optionally containing a terminal -OR''' or
- 20

~~-OC(O)R''' moiety wherein R''' is hydrogen or C₁-C₅ straight or branched chain alkyl, (f) -R' wherein R' is C₁-C₅ straight or branched chain alkyl, or (g) -R'OR''' wherein R' is C₁-C₅ straight or branched chain alkyl and R''' is hydrogen or C₁-C₅ alkyl;~~

~~G is hydrogen; and~~

- 5 ~~R₃ is (a) C₁-C₁₂ straight or branched chain alkyl, (b) -OR''', in which R''' is a straight chain or branched C₂-C₉ alkyl which may be substituted at the terminal carbon atom by a phenyl group, or (c) -(CH₂)_nOR''' wherein n is an integer of 1 to 7 and R''' is hydrogen or C₁-C₅ alkyl.~~

10 2. The compound of claim 1, wherein R₃ is a straight or branched chain C₅-C₁₂ alkyl.

3. The compound of claim 1, wherein R₃ is 1,1-dimethyl heptyl or 1,2-dimethyl heptyl.

15 4. The compound of claim 1 wherein Q is a saturated or unsaturated ring of 4 to 8 members consisting of C with at least one of N, S, and O, said ring being optionally substituted with -COR''' or -COOR''' wherein R''' is a hydrogen or C₁-C₅ straight or branched chain alkyl.

5. The compound of claim 1, wherein R₁ is -CH₂OH, -C(O)N(R'')₂, -C(O)OR'', -COOH, an amino acid, or a carboxamide.

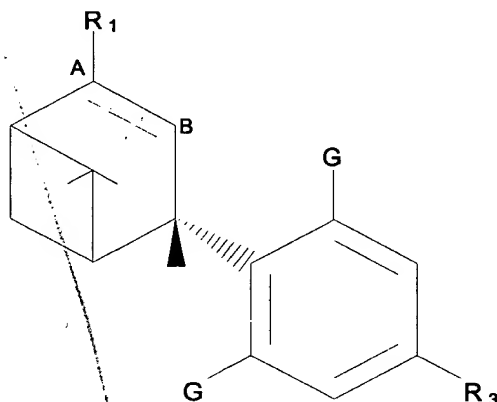
20

6. A pharmaceutical composition for treating, preventing, or managing hypertension, inflammation, peripheral pain, gastrointestinal disorders, or autoimmune diseases comprising as an active ingredient a therapeutically effective amount of a compound of claim 1.

25 7. The pharmaceutical composition of claim 6 further comprising a pharmaceutically acceptable diluent or carrier.

8. The pharmaceutical composition of claim 7, wherein the diluent is an aqueous cosolvent solution comprising a pharmaceutically acceptable cosolvent, a micellar solution or emulsion
30 prepared with natural or synthetic ionic or non-ionic surfactants, or a combination of such cosolvent and micellar or emulsion solutions.

9. A method for preventing, treating, or managing hypertension, inflammation, peripheral pain, gastrointestinal disorders, or autoimmune diseases comprising administering to an individual in need thereof a pharmaceutical composition comprising a therapeutically effective amount a compound of the general formula:



5 having the (3S,4S) configuration, and which is essentially free of the (3R,4R) enantiomer, wherein:

A---B designates an optional double bond,

R₁ is (a) -R'N(R'')₂ wherein R' is C₁-C₅ straight or branched chain alkyl and each R'', which may be the same or different, is hydrogen or C₁-C₅ straight or branched chain alkyl optionally containing a terminal -OR''' or -OC(O)R''' moiety wherein R''' is hydrogen or C₁-C₅ straight or branched chain alkyl, (b) -Q wherein Q is a heterocyclic moiety having a labile hydrogen atom so that said moiety acts as a carboxylic acid analogue, (c) -R'X wherein R' is C₁-C₅ straight or branched chain alkyl and X is halogen, (d) -R'C(O)N(R'')₂ wherein R' is a direct bond or C₁-C₅ straight or branched chain alkyl and each R'', which may be the same or different, is hydrogen or C₁-C₅ straight or branched chain alkyl optionally containing a terminal -OR''' or -OC(O)R''' moiety wherein R''' is hydrogen or C₁-C₅ straight or branched chain alkyl, (e) -R'C(O)OR'' wherein R' is a direct bond or C₁-C₅ straight or branched chain alkyl and R'' is hydrogen or C₁-C₅ straight or branched chain alkyl optionally containing a terminal -OR''' or -OC(O)R''' moiety wherein R''' is hydrogen or C₁-C₅ straight or branched chain alkyl, (f) -R' wherein R' is C₁-C₅ straight or branched chain alkyl, or (g) -R'OR''' wherein R' is C₁-C₅ straight or branched chain alkyl and R''' is hydrogen or C₁-C₅ alkyl;

G is hydrogen, halogen, or $-OR_2$ wherein R_2 is hydrogen or C_1-C_5 straight or branched chain alkyl optionally containing a terminal $-OR'''$, $-OC(O)R'''$, $C(O)OR'''$, or $-C(O)R'''$ moiety wherein R''' is hydrogen or C_1-C_5 straight or branched chain alkyl; and

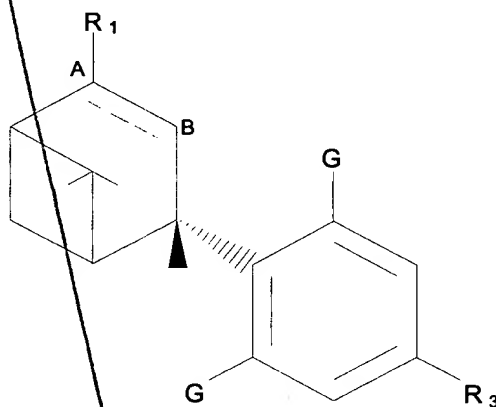
R_3 is (a) C_1-C_{12} straight or branched chain alkyl, (b) $-OR'''$, in which R''' is a straight chain or branched C_2-C_9 alkyl which may be substituted at the terminal carbon atom by a phenyl group, or (c) $-(CH_2)_nOR'''$ wherein n is an integer of 1 to 7 and R''' is hydrogen or C_1-C_5 alkyl.

10. The method of claim 9 wherein, R_1 is $-CH_2OH$, G is hydrogen or OR_2 , R_2 is a lower alkyl group, and R_3 is a straight or branched chain C_5-C_{12} alkyl.

11. The method of claim 10, wherein G is $-OCH_3$ and R_3 is 1,1-dimethyl heptyl.

12. The method of claim 10, wherein R_1 is $-CH_2OH$, G is $-OCH_3$, and R_3 is 1,1-dimethyl heptyl.

13. A method for preventing, treating, or managing tumors expressing CB2 receptors comprising administering to an individual in need thereof a pharmaceutical composition comprising a therapeutically effective amount a compound of the general formula:



having the (3S,4S) configuration, and which is essentially free of the (3R,4R) enantiomer, wherein:

A---B designates an optional double bond,

000201-12085950

R₁ is (a) -R'N(R'')₂ wherein R' is C₁-C₅ straight or branched chain alkyl and each R'', which may be the same or different, is hydrogen or C₁-C₅ straight or branched chain alkyl optionally containing a terminal -OR''' or -OC(O)R''' moiety wherein R''' is hydrogen or C₁-C₅ straight or branched chain alkyl, (b) -Q wherein Q is a heterocyclic moiety having a labile hydrogen atom so that said moiety acts as a carboxylic acid analogue, (c) -R'X wherein R' is C₁-C₅ straight or branched chain alkyl and X is halogen, (d) -R'C(O)N(R'')₂ wherein R' is a direct bond or C₁-C₅ straight or branched chain alkyl and each R'', which may be the same or different, is hydrogen or C₁-C₅ straight or branched chain alkyl optionally containing a terminal -OR''' or -OC(O)R''' moiety wherein R''' is hydrogen or C₁-C₅ straight or branched chain alkyl, (e) -R'C(O)OR'' wherein R' is a direct bond or C₁-C₅ straight or branched chain alkyl and R'' is hydrogen or C₁-C₅ straight or branched chain alkyl optionally containing a terminal -OR''' or -OC(O)R''' moiety wherein R''' is hydrogen or C₁-C₅ straight or branched chain alkyl, (f) -R' wherein R' is C₁-C₅ straight or branched chain alkyl, or (g) -R'OR''' wherein R' is C₁-C₅ straight or branched chain alkyl and R''' is hydrogen or C₁-C₅ alkyl;

15 G is hydrogen, halogen, or -OR₂ wherein R₂ is hydrogen or C₁-C₅ straight or branched chain alkyl optionally containing a terminal -OR''', -OC(O)R''', C(O)OR''', or -C(O)R'' moiety wherein R''' is hydrogen or C₁-C₅ straight or branched chain alkyl; and

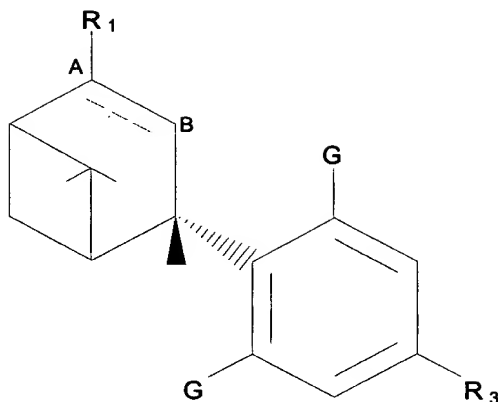
R₃ is (a) C₁-C₁₂ straight or branched chain alkyl, (b) -OR''', in which R''' is a straight chain or branched C₂-C₉ alkyl which may be substituted at the terminal carbon atom by a phenyl group, or (c) -(CH₂)_nOR''' wherein n is an integer of 1 to 7 and R''' is hydrogen or C₁-C₅ alkyl.

14. The method of claim 13 wherein, R₁ is -CH₂OH, G is hydrogen or OR₂, R₂ is a lower alkyl group, and R₃ is a straight or branched chain C₅-C₁₂ alkyl.

25 15. The method of claim 14, wherein G is -OCH₃ and R₃ is 1,1-dimethyl heptyl.

16. The method of claim 14, wherein R₁ is -CH₂OH, G is -OCH₃, and R₃ is ~~1,1-dimethyl heptyl.~~

30 17. A CB2 specific antagonist comprising a compound of the general formula:



having the (3S,4S) configuration, and which is essentially free of the (3R,4R) enantiomer, wherein:

A---B designates an optional double bond,

R₁ is (a) -R'N(R'')₂ wherein R' is C₁-C₅ straight or branched chain alkyl and each R'',
 5 which may be the same or different, is hydrogen or C₁-C₅ straight or branched chain alkyl optionally containing a terminal -OR''' or -OC(O)R''' moiety wherein R''' is hydrogen or C₁-C₅ straight or branched chain alkyl, (b) -Q wherein Q is a heterocyclic moiety having a labile hydrogen atom so that said moiety acts as a carboxylic acid analogue, (c) -R'X wherein R' is C₁-C₅ straight or branched chain alkyl and X is halogen, (d) -R'C(O)N(R'')₂ wherein R' is a direct
 10 bond or C₁-C₅ straight or branched chain alkyl and each R'', which may be the same or different, is hydrogen or C₁-C₅ straight or branched chain alkyl optionally containing a terminal -OR''' or -OC(O)R''' moiety wherein R''' is hydrogen or C₁-C₅ straight or branched chain alkyl, (e) -R'C(O)OR'' wherein R' is a direct bond or C₁-C₅ straight or branched chain alkyl and R'' is hydrogen or C₁-C₅ straight or branched chain alkyl optionally containing a terminal -OR''' or
 15 -OC(O)R''' moiety wherein R''' is hydrogen or C₁-C₅ straight or branched chain alkyl, (f) -R' wherein R' is C₁-C₅ straight or branched chain alkyl, or (g) -R'OR''' wherein R' is C₁-C₅ straight or branched chain alkyl and R''' is hydrogen or C₁-C₅ alkyl;

G is hydrogen, halogen, or -OR₂ wherein R₂ is hydrogen or C₁-C₅ straight or branched chain alkyl optionally containing a terminal -OR''', -OC(O)R''', C(O)OR''', or -C(O)R'' moiety
 20 wherein R''' is hydrogen or C₁-C₅ straight or branched chain alkyl; and

b3

add
bif

[illegible]